Please substitute the following claim set for those currently of record:

- 1. -36. (Cancelled)
- (Currently amended) A method for analyzing nucleotide sequences variations.

forming microemulsions comprising one or more species of analyte DNA molecules;

amplifying analyte DNA molecules in the microemulsions in the presence of reagent beads, wherein the reagent beads are bound to a plurality of molecules of a primer for amplifying the analyte DNA molecules, whereby product beads are formed which are bound to a plurality of copies of one species of analyte DNA molecule;

separating the product beads from analyte DNA molecules which are not bound to product beads;

determining a sequence feature of the one species of analyte DNA molecule which is bound to the product beads by flow cytometry\*

isolating using fluorescence activated cell sorting product beads which are bound to a plurality of copies of a first species of analyte DNA molecule from product beads which are bound to a plurality of copies of a second species of analyte DNA molecule.

- 38. (Cancelled)
- (Currently amended) A method for analyzing nucleotide sequences variations, comprising:

forming microemulsions comprising one or more species of analyte DNA molecules;

amplifying analyte DNA molecules in the microemulsions in the presence of reagent beads, wherein the reagent beads are bound to a plurality of molecules of a primer for amplifying the analyte DNA molecules, whereby product beads are formed which are bound to a plurality of copies of one species of analyte DNA molecule;

separating the product beads from analyte DNA molecules which are not bound to product beads:

determining a sequence feature of the one species of analyte DNA molecule which is bound to the product beads;

isolating product beads which are bound to a plurality of copies of a first the one species of analyte DNA molecule from product beads which are bound to a plurality of copies of a second species of analyte DNA molecule;

amplifying the  $\frac{\text{First}}{\text{one}}$  species of analyte DNA molecule from the isolated product beads.

- 40. (Cancelled)
- 41. (Cancelled)
- 42. (Cancelled)
- 43. (Currently amended) A method for analyzing nucleotide sequences variations, comprising:

forming microemulsions comprising one or more species of analyte DNA molecules:

amplifying analyte DNA molecules in the microemulsions in the presence of reagent beads, wherein the reagent beads are bound to a plurality of molecules of a primer for amplifying the analyte DNA molecules, whereby product beads are formed which are bound to a plurality of copies of one species of analyte DNA molecule;

separating the product beads from analyte DNA molecules which are not bound to product beads;

determining a sequence feature of the one species of analyte DNA molecule which is bound to the product beads by hybridization to oligonucleotide probes which are differentially labeled.

44. (Currently amended) A method for analyzing nucleotide sequences variations, comprising:

forming microemulsions comprising one or more than one species of analyte DNA molecules;

amplifying analyte DNA molecules in the microemulsions in the presence of reagent beads, wherein the reagent beads are bound to a plurality of molecules of a primer for amplifying the analyte DNA molecules, whereby product beads are formed which are bound to a plurality of copies of one species of analyte DNA molecule;

separating the product beads from analyte DNA molecules which are not bound to product beads;

determining relative or absolute amounts of product beads comprising one or more sequence features a first species of analyte DNA molecule as a fraction of product beads.

- (Currently amended) The method of claim 44 wherein the relative or absolute amounts are determined using flow cytometry.
- 46. -59. (Cancelled)
- 60. (Currently amended) A method for isolating nucleotide sequences variations comprising:

forming microemulsions comprising one or more than one species of analyte DNA molecules;

amplifying analyte DNA molecules in the microemulsions in the presence of reagent beads, wherein the reagent beads are bound to a plurality of molecules of a primer for amplifying the analyte DNA molecules, whereby product beads are formed which are bound to a plurality of copies of one species of analyte DNA molecule;

separating the product beads from analyte DNA molecules which are not bound to product beads;

isolating using fluorescence activated cell sorting product beads which are bound to a plurality of copies of a first species of analyte DNA molecule from product beads which are bound to a plurality of copies of a second species of analyte DNA molecule.

- 61. (Cancelled)
- 62. (Currently amended) A method for isolating nucleotide sequences variations, comprising:

forming microemulsions comprising <del>one or</del> more <u>than one</u> species of analyte DNA molecules;

amplifying analyte DNA molecules in the microemulsions in the presence of reagent beads, wherein the reagent beads are bound to a plurality of molecules of a primer for amplifying the analyte DNA molecules, whereby product beads are formed which are bound to a plurality of copies of one species of analyte DNA molecule;

separating the product beads from analyte DNA molecules which are not bound to product beads;

isolating product beads which are bound to a plurality of copies of a first species of analyte DNA molecule from product beads which are bound to a plurality of copies of a second species of analyte DNA molecule;

amplifying the first species of analyte DNA molecule from the isolated product beads.

63, -84, (Cancelled)

## 85. (New) A method for analyzing nucleotide sequences, comprising:

forming microemulsions comprising more than one species of analyte DNA molecules:

amplifying analyte DNA molecules in the microemulsions in the presence of reagent beads, wherein the reagent beads are bound to a plurality of molecules of a primer for amplifying the analyte DNA molecules, whereby product beads are formed which are bound to a plurality of copies of a first species of analyte DNA molecule and product beads are formed which are bound to a plurality of copies of a second species of analyte DNA molecules:

separating the product beads from analyte DNA molecules which are not bound to product beads;

determining proportion of product beads comprising the first species of analyte DNA molecule to product beads comprising the second species of analyte DNA molecule

- 86. (New) The method of claim 44 wherein the first species of analyte DNA molecule is a mutant allele.
- 87. (New) The method of claim 85 wherein the first species of analyte DNA molecule is a mutant allele.
- (New) The method of claim 87 wherein the second species of analyte DNA molecule is a wild-type allele.

- 89. (New) The method of claim 44 wherein the first species of analyte DNA molecule is a wild-type allele.
- 90. (New) The method of claim 85 wherein the first species of analyte DNA molecule is a wild-type allele.